AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in this application.

Please amend the claims to read as set forth below.

1. (Currently Amended) Tablet for oral administration that disintegrates quickly in the oral cavity in less than 30 seconds, comprising:

i) Spray-dried mannitol in a proportion of at least 59.5%;

ii) active ingredient in a proportion below or equal to 10%, as a fine powder in which at least 90% in weight of the active ingredient has a particle size less than 100 μm;

iii) Microcrystalline cellulose in a proportion from 10 to 18%, with an average particle size of approximately 50 μm where at least 99% in weight of microcrystalline cellulose has a particle size below 250 μm;.

iv) Sodium croscarmellose in a proportion from 1 to 4%; and

v) A lubricant agent in a proportion from 0.5 to 2% in weight,

where, unless specified otherwise, the percentages are expressed in percent weight of the total weight of the tablet.

2. (Currently Amended) Tablet for oral administration according to claim 1, ~~characterised in that~~ wherein it has a friability below 0.5% ~~according to Ph. Eur. 2.9.7~~.

3. (Currently Amended) Tablet for oral administration according to claim 2, ~~characterised in that~~ wherein it has a friability below 0.2% ~~according to Ph. Eur. 2.9.7~~.

4. (Currently Amended) Tablet for oral administration according to claim 1, ~~characterised in that~~ wherein it has an apparent density from 1.1 to 1.3 g/ml.

5. (Currently Amended) Tablet for oral administration according to claim 1, ~~characterised in that~~ wherein it has a flavouring agent in a proportion from 0.5 to 2% in weight of the total weight of the tablet.

6. (Currently Amended) Tablet for oral administration according to claim 5, ~~characterised in that~~ wherein it has an artificial sweetener in a proportion from 0.5 to 2% in weight of the total weight of the tablet.

7. (Currently Amended) Tablet for oral administration according to claim 1, ~~characterised in that~~ wherein it has a humidity adsorbing agent in a proportion from 0.1 to 0.5% in weight of the total weight of the tablet.

8. (Currently Amended) Tablet for oral administration according to claim 1, ~~characterised in that~~ wherein it has an anti-adherent agent in a proportion from 0.5 to 2% in weight of the total weight of the tablet.

9. (Currently Amended) Tablet for oral administration according to claim 1, ~~characterised in that~~ wherein the proportion of insoluble elements is below 20% in weight of the total weight of the tablet.

10. (Currently Amended) Tablet for oral administration according to ~~any of previous claims, characterised in that it has a round shape, flat, bevelled with a~~ claim 1, wherein said tablet has a round shape and is flat and bevelled, said tablet having a thickness from 1.8 to 2.2 mm.

11. (Currently Amended) Tablet for oral administration according to claim 10, ~~characterised in that~~ wherein it disintegrates quickly in the oral cavity in less than 20 seconds.

12. (Currently Amended) Process for obtaining a tablet for oral administration ~~as defined in any of claims 1 to 11, characterised in that it comprises~~ comprising the following steps:

I) Sieving and mixing the components except for the lubricant agent;

ii) Sieving the lubricant agent;

iii) Mixing of all the components; and

iv) ~~Direct~~ Directly compression of the final mixture.

13. (Currently Amended) Process for obtaining a tablet according to claim 12, ~~characterised in that~~ wherein said final mixture has a flowability below or equal to 10 seconds ~~according to Ph. Eur. 2.9.16~~.

14. (Currently Amended) Process for obtaining a tablet according to claim 12, ~~characterised in that~~ wherein said final mixture has an ability to settle below or equal to 20 ml ~~according to Ph. Eur. 2.9.15~~.